IN THE CLAIMS

- 1. (AMENDED) A method of treating a proliferative disease breast cancer, lung cancer, pancreatic cancer, colon cancer, myeloid leukemia, melanoma, thyroid follicular cancer, bladder carcinoma, glioma, myelodysplastic syndrome, ovarian cancer or prostate cancer in a patient in need of such treatment, comprising administering to said patient, a therapeutically effective amount of a combination of (1) a liposomal anthracycline composition in association with and (2) a growth factor receptor inhibitor. Trastuzumab; wherein said Trastuzumab is administered prior to, concurrently with or after the administration of said liposomal anthracycline composition; and wherein said liposomal anthracycline composition is pegylated liposomal doxorubicin comprising:
 - a) doxorubicin HCl;
- b) N-(carbonyl-methoxypolyethylene glycol 2000)-1,2-distearoyl-sn-glycero-3-phosphoethanolamine sodium salt;
 - fully hydrogenated soy phosphatidylcholine;
 - d) cholesterol;

histidine, hydrochloric acid and/or sodium hydroxide, ammonium sulfate, and sucrose; wherein the weight percentage ratio of a:b:c:d is about 1.0 :1.60 : 4.80 : 1.60 mg/mL respectively

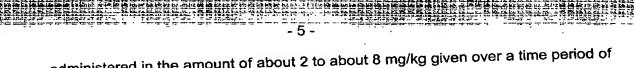
- 2. (AMENDED) The method of Claim 1, wherein said growth factor receptor inhibitor is an antibody directed against the extracellular domain of a growth factor receptor, and said patient is a treatment experienced patient having a proliferative disease and/or has at least one cardiac risk factor and/or has had previous anthracycline therapy.
- 3. (Original) The method of Claim 2, further comprising an additional antineoplastic agent.
 - 4. Cancel without prejudice.

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5. Cancel without prejudice

- - 6. Cancel without prejudice.
 - 7. Cancel without prejudice.
 - 8. (AMENDED) The method of Claim [[6]] 2 wherein the pegylated liposomal anthracycline composition and the antibody directed against the extracellular domain of a growth factor receptor Trastuzumab are administered sequentially.
 - 9. (AMENDED) The method of Claim [[6]] 2 wherein the pegylated liposomal anthracycline composition is administered first.
 - 10. (AMENDED) The method of Claim [[6]] 2 wherein the antibody directed against the extracellular domain of a growth factor receptor Trastuzumab is administered first.
 - 11. Cancel without prejudice.
 - 12. Cancel without prejudice.
 - 13. Cancel without prejudice.
 - 14. (AMENDED) The method of Claim [[11]] 3 wherein the additional antineoplastic agent is selected from the group consisting of: Uracil mustard, Cyclophosphamide, Ifosfamide, Melphalan, Chlorambucil, Temozolomide, 5-Fluorouracil, Fludarabine phosphate, Gemcitabine, Paclitaxel, Docetaxel, Interferons, Etoposide, Tamoxifen, Leuprolide, Flutamide, Toremifene, Cisplatin, Carboplatin, Navelbene, CPT-11, Anastrazole, Letrazole, and Capecitabine.
 - 15. (AMENDED) The method of Claim [[11]] 3 wherein (1) the pegylated liposomal anthracycline composition, (2) the antibody directed against the extracellular domain of a growth factor receptor Trastuzumab, and (3) the additional antineoplastic agent are administered sequentially.

- 16. (AMENDED) The method of Claim [[11]] 3 wherein the additional antineoplastic agent is Cyclophosphamide.
 - 17. Cancel without prejudice.
 - 18. Cancel without prejudice.
- 19. (AMENDED) The method of claim 4 Claim 1 wherein the pegylated liposomal anthracycline composition is administered in the amount of about 20 to about 50 mg/m², given over a time period of about 45 to about 90 minutes, every three to four weeks.
- 20. (AMENDED) The method of claim 1 wherein the antibody directed against the extracellular domain of a growth factor receptor Trastuzumab is administered first in the amount of about 2 to about 6 mg/kg given once over a time period of about 60 to about 90 minutes and subsequently administered in the amount of about 2 to about 6 mg/kg given over a time period of about 60 to 90 minutes every one to four weeks.
- 21. (AMENDED) The method of claim 5 Claim 3 wherein the additional antineoplastic agent is administered in the amount of about 400 to about 600 mg/m² given over a time period of about 20 to about 60 minutes every two to four weeks.
 - 22. Cancel without prejudice.
 - 23. Cancel without prejudice.
 - 24. (AMENDED) The method of claim 11 Claim 3 wherein
- a) the pegylated liposomal doxorubicin composition is administered in the amount of about 20 to about 50 mg/m² given over a time period of about 45 to about 90 minutes every three to four weeks [[.]] :
- b) Trastuzumab is administered first in the amount of about 2 to about 8 mg/kg given over a time period of about 60 to about 90 minutes and subsequently



administered in the amount of about 2 to about 8 mg/kg given over a time period of about 60 to about 90 minutes every one to four weeks; and

- c) the additional antineoplastic agent is Cyclophosphamide and is administered in the amount of about 400 to about 600 mg/m² given over a time period of about 20 to about 60 minutes every two to four weeks.
- 25. (AMENDED) The method of claim Claim 24 wherein (1) the pegylated liposomal doxorubicin composition is administered first, followed by (2) Cyclophosphamide and then (3) Trastuzumab.

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